CLAIMS:

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- 1. A method for labeling a cell, the method comprising contacting the cell ex vivo with a fluorocarbon imaging reagent under conditions such that the fluorocarbon imaging reagent becomes associated with the cell.
- 5 2. The method of claim 1, wherein the fluorocarbon imaging reagent is a perfluoropolyether.
 - 3. The method of claim 1, wherein the cell is contacted with the fluorocarbon imaging reagent in the presence of an uptake enhancing reagent.
 - 4. The method of claim 3, wherein the uptake enhancing reagent comprises a cationic lipid.
 - 5. The method of claim 1, wherein at least a portion of the fluorocarbon imaging reagent is internalized into the cell.
 - 6. The method of claim 1, wherein at least a portion of the fluorocarbon imaging reagent is associated with the extracellular surface of the cell.
- 7. The method of claim 1, wherein the fluorocarbon imaging reagent is conjugated to a cellular targeting moiety.
 - 8. The method of claim 7, wherein the cellular targeting moiety comprises an antibody that binds to an epitope that is exposed to the extracellular milieu.
 - 9. The method of claim 1, wherein the fluorocarbon imaging reagent is conjugated to an internalization moiety.
 - 10. The method of claim 1, wherein the cell is a mammalian cell.
 - 11. The method of claim 1, wherein the cell is a cell of the immune system.
 - 12. The method of claim 1, wherein the cell is a dendritic cell.
 - 13. The method of claim 1, wherein the fluorocarbon imaging reagent is formulated as an emulsion.
 - 14. The method of claim 1, wherein the emulsion comprises particles having a mean diameter of between 30 and 500 nm.

15. The method of claim 1, wherein the fluorocarbon imaging reagent is a perfluoro-crown ether.

- 16. The method of claim 15, wherein the imaging reagent is a perfluro-15-crown-5-ether.
- 5 17. The method of claim 1, wherein the fluorocarbon is a perfluorinated polyether having an average formula:

$$XO(Y-O)_nZ$$

wherein Y is selected from the group consisting of:

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wherein n is an integer from 8 to 20; wherein X and Z are the same and are selected from the group consisting of perfluoroalkyls, perfluoroathers, fluoroalkyls terminated with fluoroacyl, carboxyl, amide or ester, methylols, acid chlorides, amides, amidines, acrylates and esters.

- 18. The method of claim 1, wherein the imaging reagent comprises an additional functional moiety.
 - 19. The method of claim 18, wherein the additional functional moiety is a detection moiety.
 - 20. The method of claim 19, wherein the detection moiety is selected from the group consisting of: a fluorescent detection moiety and a PET detection moiety.
 - 21. An imaging reagent having an average formula:

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$$XO(Y-O)_nZ$$

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wherein Y is selected from the group consisting of:

wherein n is an integer from 8 to 20; wherein X and Z are the same and are selected from the group consisting of perfluoroalkyls, perfluoroathers, fluoroalkyls terminated with fluoroacyl, carboxyl, amide or ester, methylols, acid chlorides, amides, amidines, acrylates and esters.

- 22. The imaging reagent of claim 21, wherein n=11.
- 23. The imaging reagent of claim 21, wherein X and Z are perfluoroethers terminated with a carboxyl group.
 - 24. The imaging reagent of claim 21, wherein each carboxyl is derivatized with a polyethylene glycol.
 - 25. The imaging reagent of claim 21, wherein X and Z are derivatized with a fluorescent detection moiety.
 - 26. A linear fluorocarbon derivatized at one or more polymer ends with at least one functional moiety, wherein the at least one functional moiety is selected from the group consisting of: a detection moiety, a hydrophilic moiety, a targeting moiety and a cellular uptake moiety.
- 27. The linear fluorocarbon of claim 26, wherein the linear fluorocarbon is a linear perfluoropolyether.
 - 28. The linear fluorocarbon of claim 26, wherein the at least one functional moiety is a detection moiety.

29. The linear fluorocarbon of claim 28, wherein the detection moiety is selected from the group consisting of: a fluorescent detection moiety and a PET detection moiety.

- 30. An emulsion comprising a perfluoropolyether and having a particle size ranging from 10 to 500 nm.
- 31. The emulsion of claim 30, wherein the emulsion is stable at temperatures ranging from 4°C to 37°C.
- 32. A method for detecting a cell in a subject, the method comprising:

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- a. administering to the subject a cell that is labeled with a fluorocarbon imaging reagent; and
- b. examining at least a portion of the subject by a nuclear magnetic resonance technique, thereby detecting a labeled cell in the subject.
- 33. The method of claim 32, wherein examining by a nuclear magnetic resonance technique comprises collecting an ¹⁹F data set.
- 15 34. The method of claim 33, further comprising collecting an ¹H data set.
 - 35. The method of claim 34, further comprising generating and comparing a ¹⁹F image and a ¹H image.
 - 36. The method of claim 32, wherein the nuclear magnetic resonance technique is magnetic resonance imaging (MRI).
- 37. The method of claim 32, wherein the nuclear magnetic resonance technique is magnetic resonance spectroscopy (MRS).
 - 38. The method of claim 32, wherein the fluorocarbon imaging reagent is a perfluoropolyether.
- 39. The method of claim 38, wherein the fluorocarbon imaging reagent is selected from the group consisting of: a linear perfluoropolyether, a cyclic perfluoropolyether and a mixture thereof.
 - 40. The method of claim 32, wherein the cell is a mammalian cell.
 - 41. The method of claim 32, wherein the cell is a cell of the immune system.

- 42. The method of claim 32, wherein the cell is a dendritic cell.
- 43. The method of claim 32, wherein the cell is administered to the subject as part of a cellular therapeutic regimen.
- 44. The method of claim 31, wherein the cell is a stem cell.
- 5 45. A labeled cellular formulation for administration to a subject, the formulation comprising:
 - a. a cell; and

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- b. a fluorocarbon imaging reagent that is associated with the cell.
- 46. The formulation of claim 45, further comprising a pharmaceutically acceptable excipient.
- 47. The formulation of claim 45, wherein the fluorocarbon imaging reagent is a perfluoropolyether.
- 48. The formulation of claim 45, wherein at least a portion of the fluorocarbon imaging reagent is internalized into the cell.
- 49. The formulation of claim 45, wherein at least a portion of the fluorocarbon imaging reagent is associated with the extracellular surface of the cell.
 - 50. The formulation of claim 45, wherein the fluorocarbon imaging reagent is conjugated to a cellular targeting moiety.
 - 51. The formulation of claim 50, wherein the cellular targeting moiety comprises an antibody that binds to an epitope that is exposed to the extracellular milieu.
 - 52. The formulation of claim 45, wherein the fluorocarbon imaging reagent is conjugated to an internalization moiety.
 - 53. The formulation of claim 45, wherein the cell is a mammalian cell.
- 25 54. The formulation of claim 45, wherein the cell is a cell of the immune system.
 - 55. The formulation of claim 45, wherein the cell is a dendritic cell.
 - 56. The formulation of claim 45, wherein the cell is prepared for use in a cellular therapeutic regimen.

- 57. A method for detecting transplanted cells in a transplant recipient, the method comprising:
 - a. administering cells for transplant to a transplant recipient, at least a
 portion of which cells for transplant are labeled with a fluorocarbon
 imaging reagent;
 - examining at least a portion of the subject by a nuclear magnetic resonance technique, thereby detecting the labeled cells.
- 58. The method of claim 57, wherein the location and optionally the trafficking of labeled cells is detected in the transplant recipient.
- 59. The method of claim 56, wherein the nuclear magnetic resonance technique is selected from the group consisting of: magnetic resonance imaging and magnetic resonance spectroscopy.
 - 60. The method of claim 56, wherein the transplant recipient is a bone marrow transplant recipient.
- 15 61. The method of claim 56, wherein the cells for transplant comprise hematopoietic stem cells.

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- 62. The method of claim 56, wherein the cells for transplant are derived from bone marrow, cord blood or peripheral blood.
- 63. The method of claim 56, wherein the transplant recipient is the recipient of a donor organ.
 - 64. The method of claim 62, wherein at least a portion of the cells of the donor organ are labeled with a fluorocarbon imaging reagent.